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In the Claims

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The listing of claims will replace all prior versions and listings of claims in the application.

Listings of claims

- 1. (Original) A method for identifying compounds that modulate topoisomerase activity, said method comprising:
- a) providing cells expressing topoisomerase and containing a promoter sensitive to changes in DNA topology having a reporter gene operatively linked thereto;
- b) measuring the expression of said reporter gene in the presence and in the absence of a test compound;
- c) comparing the expression of said reporter gene in the presence of said compound with the expression in the absence of said compound; and
- d) identifying a compound that modulates topoisomerase activity as one that yields an alteration in reporter gene expression in the presence of the compound relative to expression in the absence of the compound.
- 2. (Currently Amended) <u>The method of claim 1, wherein modulation of topoisomerase activity is inhibition of topoisomerase activity.</u> A method-for identifying compounds that inhibit topoisomerase activity, said-mothod comprising:
- a)——providing—cells—expressing_topoisomerase—and_containing_a—promotor—sensitive_to changes in DNA-topology-having-a-reporter-gene operatively-linked thereto;
- b) measuring the expression of said reporter gene in the presence and in the absence of a-test-compound;
- c) comparing the expression of said reporter gene in the presence of said compound with the expression in the absence of said compound; and
- d) identifying a compound that inhibits topoisomerase activity as one that yields an alteration in reporter gene expression in the presence of the compound relative to expression in the absence of the compound.
- 3. (Currently Amended) The method of claim 1-er-2, wherein the topoisomerase is a type II topoisomerase.
- 4. (Currently Amended) The method of claim 1-or-2, wherein the topoisomerase is a DNA gyrase.

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- 5. (Currently Amended) The method of claim 1—er—2, wherein the topoisomerase is a recombinant topoisomerase.
- 6. (Currently Amended) The method of claim 1-er-2, wherein the topoisomerase is a prokaryotic, eukaryotic, or viral topoisomerase.
- 7. (Currently Amended) The method of claim 1—or 2, wherein the promoter is selected from gyrA, gyrB, proU, tppB, or pC, ompF, topA, dnaA, hisD, recF, katE, katG, sodA, sodB, tonB, and laclq mutant, SEQ ID NO:1, SEQ ID NO:2, SEQ ID NO:3, SEQ ID NO:4, SEQ ID NO:5, and any functional fragment thereof.
- 8. / (Currently Amended) The method of claim 1 $\frac{1}{2}$, wherein the cells are bacterial cells.
- 9. (Original) The method of claim 8, wherein the cells are Gram-positive bacterial cells.
- 10. (Original) The method of claim 8, wherein the cells are Gram-negative bacterial cells.
- 11. (Original) The method of claim 8, wherein the cells are selected from Haemophilus influenzae, Moraxella catarrhalis, Pseudomonas aeruginosa, Escherichia coli, Chlamydia spp, Legionella spp, Staphylococcus aureus, Staphylococcus saprophyticus, Streptococcus pneumoniae, Streptococcus pyogenes, Streptococcus mutans, Enterococcus faecalis, Enterococcus faecium, Mycoplasma spp, Bacteroides spp and Clostridium spp.
- 12. (Currently Amended) The method of claim 1—or 2, wherein the cells are eukaryotic cells selected from mammalian or fungal cells.
- 13. (Original) The method of claim 12, wherein the cells are human cells, Saccharomyces spp, Aspergillus spp, or Candida spp cells.
- 14. (Currently Amended) The method of claim 1 or 2, wherein the promoter and reporter gene are provided on a plasmid.

15.-17. (Canceled)

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19. (Currently Amended) The method of claim 1-or 2, wherein the reporter gene is selected from *lacZ*, *luxABCDE* operon, *lucFF* operon, *luxAB* operon, *uidA*, *gfp*, *phoA*, *kan*, *cam*, and genes encoding reef coral fluorescent proteins.

20.-22. (Canceled)

- 23. (Original) A method for identifying compounds that modulate DNA gyrase activity, said method comprising:
- a) providing cells expressing DNA gyrase and containing a promoter sensitive to changes in DNA topology having a reporter gene operatively linked thereto;
- b) measuring the expression of said reporter gene in the presence and in the absence of a test compound; and
- c) identifying a compound that modulates DNA gyrase activity as one that yields an alteration in reporter gene expression in the presence of the compound relative to expression in the absence of the compound.
- 24. (Currently Amended) <u>The method of claim 23, wherein the modulation is inhibition of DNA gyrase activity. A-method for identifying compounds that inhibit DNA gyrase activity, eaid method-comprising:</u>
- a)——providing—cells—expressing—DNA—gyrase—and—containing—a—promoter—sensitive—to changes in-DNA-topology-having-a-reporter-gene-operatively linked-thereto;
- b)——measuring the expression of said reporter gene in the presence and in the absence of a test-compound; and
- c) identifying a compound that inhibits DNA gyrase activity as one that yields—an alteration in reporter gene expression in the presence of the compound relative to expression in the absence of the compound.
- 25. (Currently Amended) The method of claim 23-or-24, wherein the DNA gyrase is a recombinant DNA gyrase.
- 26. (Currently Amended) The method of claim 23—or 24, wherein the DNA gyrase is selected from Haemophilus influenzae, Moraxella catarrhalis, Pseudomonas aeruginosa, Escherichia coli, Chlamydia spp, Legionella spp, Staphylococcus aureus, Staphylococcus saprophyticus, Streptococcus pneumoniae, Streptococcus pyogenes, Streptococcus mutans,

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Enterococcus faecalis, Enterococcus faecium, Mycoplasma spp, Bacteroides spp and Clostridium spp.

27.-45. (Cancelled)